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IN THE CLAIMS:

Please cancel claims 3-10 without prejudice or disclaimer and add the following new claims 11-18.

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11. A compound according to claim 1 wherein R¹ and R^{1a} are independently methoxy, amino(C₃₋₅)alkyloxy, guanidino(C₃₋₅)alkyloxy, piperidyl(C₃₋₅)alkyloxy, nitro or fluoro.
12. A compound according to claim 1 wherein R³ is hydrogen; optionally substituted aminocarbonyl; optionally substituted (C₁₋₆)alkyl; carboxy(C₁₋₄)alkyl; optionally substituted aminocarbonyl(C₁₋₄)alkyl; cyano(C₁₋₄)alkyl; optionally substituted 2-oxo-oxazolidinyl or optionally substituted 2-oxo-oxazolidinyl(C₁₋₄alkyl).
13. A compound according to claim 1 wherein R³ is in the 3-position and the substituents at the 3- and 4-position of the piperidine ring are *cis*.
14. A compound according to claim 1 wherein A is NH and B is CO, or A is CHO_H and B is CH₂.
15. A compound according to claim 1 wherein R¹¹ is hydrogen.
16. A compound according to claim 1 wherein R⁴ is (C₅₋₁₂)alkyl, optionally substituted phenyl(C₂₋₃)alkyl or optionally substituted phenyl(C₃₋₄)alkenyl.
17. A pharmaceutical composition comprising a compound of formula (I) as defined in claim 1, or a pharmaceutically acceptable derivative thereof, and a pharmaceutically acceptable carrier.
18. A method of treatment of bacterial infections in mammals which method comprises the administration to a mammal in need of such treatment of an effective amount of a compound of formula (I) as defined in claim 1, or a pharmaceutically acceptable derivative thereof.